Amendments to the Claims:

This listing of claims will replace all prior versions and listings of claims in this application.

Listing of the Claims:

Claims 1-4 (cancelled).

Claim 5 (currently amended): A compound of the formula IIb:

$$R^{2d}$$
 M
 N
 $(R^{2c})_{nc}$
 R^{2a}
 N
 N
 H

$$R^{2d}$$
 M
 N
 R^{2c}
 R^{2d}
 N
 R^{2b}
 N
 H
 N
 H
 N
 H
 N
 H
 N
 H
 N
 H

wherein:

M is -CH- or -N-:

nc is 0, 1 or 2:

 $R^{2c} \ is \ linked \ to \ a \ carbon \ atom \ of \ the \ 5-membered \ ring \ and \ is \ selected \ from \ hydrogen \ and \ methyl;$

$$\begin{split} R^{2d} & \text{ is linked to a carbon atom of the 6-membered ring and is selected from hydrogen and fluoro;} \\ R^{2a} & \text{ and } R^{2b} & \text{ are each independently selected from hydrogen, hydroxy, halogeno, cyano, nitro,} \\ & \text{ trifluoromethyl, } C_{1-3} & \text{alkyl, } C_{1-3} & \text{alkoxy, } C_{1-3} & \text{alkylsulphanyl, -NR}^{3a} & \text{R}^{4a} & \text{ (wherein } R^{3a} & \text{and } R^{4a}, \\ & \text{ which may be the same or different, each represents hydrogen or } C_{1-3} & \text{alkyl), and } Q^t X^t & Q^t X^t & \text{ wherein } Q^t & \text{ is selected from one of the following groups:} \end{split}$$

 C₁₋₄alkyl-Q¹³-C(O)-C₁₋₄alkyl-Q¹⁴ wherein Q¹³ and Q¹⁴ are each independently selected from pyrrolidinyl, piperidinyl, piperazinyl,



wherein Q14 is linked to C1-4alkanoyl through a nitrogen atom;

 Q² (wherein Q² is a 5-6-membered heterocyclic group selected from pyrrolidinyl, piperidinyl, piperazinyl,



which heterocyclic group bears either one substituent selected from methylenedioxy or ethylenedioxy to form a bicyclic ring, or bears at least one substituent selected from C24 alkanoylC1-3 alkyl and optionally bears a further 1 or 2 substituents selected from C25 alkenyl, C2-5 alkynyl, C1-6 fluoroalkyl, C1-6 alkanoyl, C2-4 alkanoylC1-3 alkyl, aminoC15 alkanoyl, C1-4 alkylaminoC1-6 alkanoyl, di(C1-4 alkyl)aminoC1-6 alkanoyl, C16 fluoroalkanoyl, carbamoyl, C1-4 alkylcarbamoyl, di(C1-4 alkyl)carbamoyl, carbamoylC16 alkyl, C1-4 alkylcarbamoylC1-6 alkyl, di(C1-4 alkyl)carbamoylC1-6 alkyl, C1-6 fluoroalkylsulphonyl, oxo, hydroxy, halogeno, cyano, C1-4 cyanoalkyl, C1-4 alkyl, C14 alkoxycarbonyl, C1-4 alkoxy, C1-4 alkylamino, di(C1-4 alkyl)amino, C1-4 alkyl, C14 alkyl, di(C1-4 alkyl)aminoC1-4 alkyl, C1-4 alkylaminoC1-4 alkyl, di(C1-4 alkyl)aminoC1-4 alkyl, di(C1-4 alkyl)aminoC1-4 alkoxy, di(C1-4 alkyl)aminoC1-4 alkoxy and a group -(-O-)(C1-4 alkyl)aringD (wherein f is 0 or 1, g is 0 or 1 and ring D is a 5-6-membered saturated or partially unsaturated heterocyclic group with 1-2

heteroatoms, selected independently from O, S and N, which cyclic group may bear one or more substituents selected from $C_{1:4}$ alkyl)); and

3) C₁₋₅alkylQ² (wherein Q² is as defined herein);

and X1 is O;

and additionally wherein any $C_{1.5}$ alkyl group in Q^1X^1 - which is linked to X^1 may bear one or more substituents selected from hydroxy, halogeno and amino;

Za is -O- or -S-;

with the proviso that at least one of R^{2a} and R^{2b} is $Q^tX^t - Q^tX^t - w$ wherein Q^t and X^t are as defined herein;

or a pharmaceutically-acceptable salt thereof.

Claim 6 (currently amended): The compound according to claim 5 wherein one of R^{2a} and R^{2b} is methoxy and the other is $Q^{t}X^{t}Q^{t}X^{l}$ wherein X^{t} and Q^{t} are as defined in claim 5.

Claim 7 (currently amended): The compound according to claim 5 wherein one of R^{2a} and R^{2b} is methoxy and the other is $Q^{4}X^{4}Q^{1}X^{1}$ wherein X^{1} is -O- and Q^{1} is

C₁₋₄alkyl-Q¹³-C(O)-C₁₋₄alkyl-Q¹⁴ wherein Q¹³ and Q¹⁴ are each independently selected from pyrrolidinyl, piperidinyl, piperazinyl,

wherein Q14 is linked to C1-6alkanoyl through a nitrogen atom.

Claim 8 (currently amended): The compound according to claim 5 wherein one of R^{2a} and R^{2b} is methoxy and the other is Q^4X^4 Q^4X^4 wherein X^1 is -O- and Q^1 is selected from one of the following groups:

 Q² (wherein Q² is a 5-6-membered heterocyclic group selected from pyrrolidinyl, piperidinyl, piperazinyl,

- which heterocyclic group bears either one substituent selected from methylenedioxy or ethylenedioxy to form a bicyclic ring, or bears one substituent selected from C₂₋₄alkanoylC₁₋₃alkyl; and
- C₁₋₅alkylQ² (wherein Q² is as defined herein).

Claim 9 (previously presented): The compound according to claim 7 or claim 8 wherein \mathbb{R}^{2a} is methoxy.

Claim 10 (previously presented): The compound according to claim 5 selected from:

- 7-{[1-(acetylmethyl)piperidin-4-yl]methoxy}-6-methoxy-4-[(3-methyl-1H-indol-5-yl)oxy]quinazoline,
- 7-{[1-(acetylmethyl)piperidin-4-yl]methoxy}-6-methoxy-4-[(2-methyl-1H-indol-5-yl)oxy]quinazoline,
- 6-methoxy-4-[(3-methyl-1H-indol-5-yl)oxy]-7-{[1-(pyrrolidin-1-ylacetyl)piperidin-4-yl]methoxy}quinazoline,
- 6-methoxy-4-[(2-methyl-1*H*-indol-5-yl)oxy]-7-{[1-(pyrrolidin-1-ylacetyl)piperidin-4-yl]methoxy}quinazoline,
- 6-methoxy-4-[(2-methyl-1*H*-indol-5-yl)oxy]-7-[2-(tetrahydro-5*H*-[1,3]dioxolo[4,5-c]pyrrol-5-yl)ethoxy]quinazoline,
- 6-methoxy-4-[(3-methyl-1H-indol-5-yl)oxy]-7-[2-(tetrahydro-5H-[1,3]dioxolo[4,5-c]pyrrol-5-yl)ethoxy]quinazoline,
- 4-[(2,3-dimethyl-1*H*-indol-5-yl)oxy]-6-methoxy-7-[2-(tetrahydro-5*H*-[1,3]dioxolo[4,5-c]pyrrol-5-yl)ethoxy]quinazoline,
- 4-[(4-fluoro-2-methyl-1*H*-indol-5-yl)oxy]-6-methoxy-7-[2-(tetrahydro-5*H*-[1,3]dioxolo[4,5-c]pyrrol-5-yl)ethoxy]quinazoline,
- 7-{2-[4-(acetylmethyl)piperazin-1-yl]ethoxy}-4-[(2,3-dimethyl-1H-indol-5-yl)oxy]-6-methoxyquinazoline,
- 7-{2-[4-(acetylmethyl)piperazin-1-yl]ethoxy}-6-methoxy-4-[(3-methyl-1H-indol-5-yl)oxy]quinazoline,
- 7-{2-[4-(acetylmethyl)piperazin-1-yl]ethoxy}-6-methoxy-4-[(2-methyl-1H-indol-5-

Attorney Docket No.: 056291-5233

Page 7

yl)oxy]quinazoline,

- 7-{2-[4-(acetylmethyl)piperazin-1-yl]ethoxy}-4-[(4-fluoro-2-methyl-1H-indol-5-yl)oxy]-6-methoxyquinazoline,
- 6-methoxy-4-[(2-methyl-1*H*-indol-5-yl)oxy]-7-{2-[4-(pyrrolidin-1-ylacetyl)piperazin-1-yl]ethoxy}quinazoline,
- 7-{[1-(acetylmethyl)piperidin-4-yl]oxy}-6-methoxy-4-[(2-methyl-1H-indol-5-yl)oxy]quinazoline, and
- 7-{[1-(acctylmethyl)piperidin-4-yl]oxy}-4-[(4-fluoro-2-methyl-1*H*-indol-5-yl)oxy]-6-methoxyquinazoline,

and pharmaceutically-acceptable salts thereof.

Claims 11 - 13 (cancelled).

Claim 14 (previously presented): A pharmaceutical composition which comprises a compound of the formula IIb as defined in claim 5 or a pharmaceutically acceptable salt thereof, in association with a pharmaceutically acceptable excipient or carrier.

Claims 15-16 (cancelled).

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